EXHIBIT N

IN THE UNITED STATES DISTRICT COURT
FOR THE SOUTHERN DISTRICT OF WEST VIRGINIA
CHARLESTON DIVISION

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IN RE: ETHICON, INC. PELVIC REPAIR SYSTEM PRODUCTS LIABILITY LITIGATION : MDL NO. 2327

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THIS DOCUMENT RELATES TO ALL CASES

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Thursday, June 13, 2013

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CONFIDENTIAL - SUBJECT TO PROTECTIVE ORDER

Videotaped Deposition of JENNIFER M. PAINE held at Riker Danzig Scherer Hyland Perretti LLP, Headquarters Plaza, One Speedwell Avenue, Morristown, New Jersey, on the above date, beginning at 9:38 a.m., before Kimberly A. Overwise, a Certified Realtime Reporter, Certified Court Reporter, and Notary Public.

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It looks like that's about a two-year job; correct?

A That's correct.

Q Did you have any involvement with TVT mesh in the second job as manager of regulatory affairs from July of 2005 to July of 2007?

A It's possible that I had some involvement with TVT in that time frame. It's difficult for me to say exactly when I would have. What I can tell you is that the early part of that time frame I believe I would have only been working on new product development projects, so not necessarily covering products that were already launched in the market at that time.

Q Okay. So if we then go from July of 2007 to December of 2008, it says worldwide director of regulatory affairs; correct?

A That's correct.

Q During that time frame, were you involved with TVT mesh?

A At that time I believe that I was covering the Women's Health & Urology business in total and so, yes, I would have had oversight to folks who were working on the TVT product line.

Q Okay. Would you say most of your involvement with TVT mesh would have been during

that time period of July 2007 to December of 2008?

A Probably the majority of it would be. There may have been some, as I said, prior to that.

Q So it looks like that's a period of about 17 months; correct?

A Yes.

Q Okay. And during your time period -- what does a worldwide director of regulatory affairs for women's health do?

A So in that role I was a participant on the Women's Health & Urology board. Again, I'm not exactly sure of what time frame that role was present. But I had oversight to the regulatory team that was supporting the -- all of the women's health products, which was a fairly expansive portfolio of products, including some hardware and software devices as well as the mesh products.

Q Okay. What do you do I mean on --

A Oh, I'm sorry.

Q On a day-to-day basis, I mean, what does the worldwide director of regulatory affairs do? Do you walk into the office, read catalogs, sip coffee, go talk to the secretaries? You know, I mean, what do you do?

MS. KABBASH: Objection.